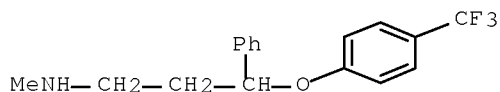


L1 1 S US 20060241102/PN

FILE 'REGISTRY' ENTERED AT 08:41:48 ON 15 OCT 2009
L2 1 S 9001-66-5/RN
SET NOTICE 1 DISPLAY
SET NOTICE LOGIN DISPLAY

FILE 'REGISTRY' ENTERED AT 08:42:02 ON 15 OCT 2009
L3 1 S 56296-78-7/RN



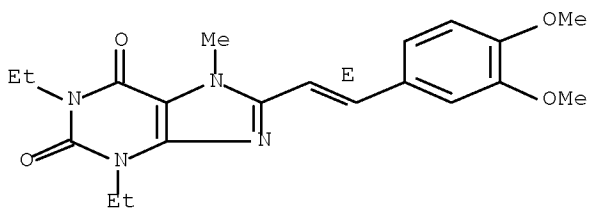
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SET NOTICE 1 DISPLAY
SET NOTICE LOGIN DISPLAY

FILE 'REGISTRY' ENTERED AT 08:42:26 ON 15 OCT 2009
E KW6002/CN
SET EXPAND CONTINUOUS
L4 1 S KW 6002/CN
E KW 6002/CN
L5 1 S E15

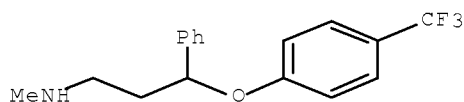
L5 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2009 ACS on STN
RN 155270-99-8 REGISTRY
ED Entered STN: 24 May 1994
CN 1H-Purine-2,6-dione, 8-[(1E)-2-(3,4-dimethoxyphenyl)ethenyl]-1,3-diethyl-
3,7-dihydro-7-methyl- (CA INDEX NAME)
OTHER CA INDEX NAMES:
CN 1H-Purine-2,6-dione, 8-[2-(3,4-dimethoxyphenyl)ethenyl]-1,3-diethyl-3,7-
dihydro-7-methyl-, (E)-
OTHER NAMES:
CN Istradefylline
CN KW 6002
FS STEREOSEARCH
MF C20 H24 N4 O4
CI COM
SR CA
LC STN Files: ADISINSIGHT, ADISNEWS, AGRICOLA, BIOSIS, BIOTECHNO, CA,
CAPLUS, CASREACT, CBNB, CHEMCATS, CIN, EMBASE, IMSPATENTS, IMSRESEARCH,
IPA, MEDLINE, MRCK*, PHAR, PROMT, PROUSDDR, RTECS*, SYNTHLINE, TOXCENTER, USAN, USPAT2, USPATFULL
(*File contains numerically searchable property data)

Double bond geometry as shown.



E FLUOXETINE/CN
 L6 1 S E27

 L6 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2009 ACS on STN
 RN 54910-89-3 REGISTRY
 ED Entered STN: 16 Nov 1984
 CN Benzenepropanamine, N-methyl-γ-[4-(trifluoromethyl)phenoxy]- (CA INDEX NAME)
 OTHER CA INDEX NAMES:
 CN Benzenepropanamine, N-methyl-γ-[4-(trifluoromethyl)phenoxy]-, (±)-
 OTHER NAMES:
 CN (±)-Fluoxetine
 CN (±)-N-Methyl-3-phenyl-3-[4-(trifluoromethyl)phenoxy]propylamine
 CN 3-(p-Trifluoromethylphenoxy)-N-methyl-3-phenylpropylamine
 CN Deprex
 CN dl-3-(p-Trifluoromethylphenoxy)-N-methyl-3-phenylpropylamine
 CN Fluoxetin Ratiopharm
 CN Fluoxetine
 CN Fluoxin
 CN Fluval
 CN N-Methyl-3-(p-trifluoromethylphenoxy)-3-phenylpropylamine
 CN Nikomed
 CN NSC 283480
 CN Symbiax
 DR 57226-07-0, 52341-67-0
 MF C17 H18 F3 N O
 CI COM
 LC STN Files: ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, AQUIRE, BEILSTEIN*, BIOSIS, BIOTECHNO, CA, CABA, CAPLUS, CASREACT, CBNB, CHEMCATS, CHEMINFORMRX, CHEMLIST, CIN, CSCHEM, CSNB, DDFU, DRUGU, EMBASE, HSDB*, IFICDB, IFIPAT, IFIUDB, IMSDRUGNEWS, IMSPATENTS, IMSPRODUCT, IMSRESEARCH, IPA, MEDLINE, MRCK*, MSDS-OHS, PHAR, PROMT, PROUSDDR, PS, RTECS*, SPECINFO, SYNTHLINE, TOXCENTER, USAN, USPAT2, USPATFULL, VETU
 (*File contains numerically searchable property data)
 Other Sources: WHO



FILE 'HCAPLUS' ENTERED AT 08:43:33 ON 15 OCT 2009

FILE 'HCAPLUS' ENTERED AT 08:44:03 ON 15 OCT 2009

L7 595 S L3
L8 106 S L5
L9 5344 S L6
L10 2 S L7 AND L8

L10 ANSWER 1 OF 2 HCAPLUS COPYRIGHT 2009 ACS on STN

TI Methods and compositions for the treatment of neurodegenerative disorders

ACCESSION NUMBER: 2008:1337969 HCAPLUS Full-text

DOCUMENT NUMBER: 149:525459

TITLE: Methods and compositions for the treatment of neurodegenerative disorders

INVENTOR(S): Jin, Xiaowei; Staunton, Jane; Macdonald, Douglas;

Dong, Hualing; Kifle, Lydia

PATENT ASSIGNEE(S): Combinatorx, Incorporated, USA; CHDI, Inc.

SOURCE: PCT Int. Appl., 123pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2008133884	A2	20081106	WO 2008-US5194	
20080423				
W:	AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW			
RW:	AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, NO, PL, PT, RO, SE, SI, SK,			

TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE,
 SN, TD,
 TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG,
 ZM, ZW,
 AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
 PRIORITY APPLN. INFO.: US 2007-925753P P
 20070423 US 2007-958774P P
 20070709

L10 ANSWER 2 OF 2 HCAPLUS COPYRIGHT 2009 ACS on STN
 TI Medicinal compositions containing adenosine A2A receptor
 antagonists and

other antidepressants
 ACCESSION NUMBER: 2005:99358 HCAPLUS Full-text
 DOCUMENT NUMBER: 142:162694
 TITLE: Medicinal compositions containing adenosine
 A2A
 receptor antagonists and other antidepressants
 INVENTOR(S): Kase, Hiroshi; Kobayashi, Minoru; Shiozaki,
 Shizuo;
 Mori, Akihisa; Seno, Naoki
 PATENT ASSIGNEE(S): Kyowa Hakko Kogyo Co., Ltd., Japan
 SOURCE: PCT Int. Appl., 47 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: Japanese
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005009444	A1	20050203	WO 2004-JP10758	
20040722				
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CA, CH,	CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI,			
GB, GD,	GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR,			
KZ, LC,	LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ,			
NA, NI,	NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK,			
SL, SY,	TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA,			
ZM, ZW				
RW:	BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM,			
ZW, AM,	AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ,			
DE, DK,	EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT,			
RO, SE,	SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML,			
MR, NE,	SN, TD, TG			

CA 2533117 A1 20050203 CA 2004-2533117
 20040722
 EP 1655029 A1 20060510 EP 2004-748023
 20040722
 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE,
 MC, PT,
 IE, SI, FI, RO, CY, TR, BG, CZ, EE, HU, PL, SK
 US 20060241102 A1 20061026 US 2006-565239
 20060119
 NO 2006000958 A 20060425 NO 2006-958
 20060227
 PRIORITY APPLN. INFO.: JP 2003-201549 A
 20030725 WO 2004-JP10758 W
 20040722
 IC ICM A61K031-52
 ICS A61K031-137; A61K031-335; A61K031-343; A61K031-36; A61K031-
 38;
 A61K031-381; A61K031-435; A61K031-496; A61K031-5375; A61K031-
 55;
 A61K031-553; A61P025-24
 CC 63-6 (Pharmaceuticals)
 Section cross-reference(s): 1
 IT ~~56296-78-7~~, Fluoxetine hydrochloride 99300-78-4, Venlafaxine
 hydrochloride 155270-99-8
 RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL
 (Biological study); USES (Uses)
 (medicinal compns. containing adenosine A2A receptor
 antagonists and other
 antidepressants)

 L11 5 S L8 AND L9
 L12 4 S L11 NOT L10
 L13 1 S L12 AND (PY<2003 OR AY<2003 OR PRY<2003)

 L13 ANSWER 1 OF 1 HCAPLUS COPYRIGHT 2009 ACS on STN
 TI Combination therapy including cyclooxygenase 2 (COX2) inhibitor(s)
 for the
 treatment of Parkinson's disease
 ACCESSION NUMBER: 2003:855794 HCAPLUS Full-text
 DOCUMENT NUMBER: 139:345938
 TITLE: Combination therapy including cyclooxygenase 2
 (COX2)
 inhibitor(s) for the treatment of Parkinson's
 disease
 INVENTOR(S): Stephenson, Diane T.; Isakson, Peter C.;
 Maziasz,
 Timothy J.
 PATENT ASSIGNEE(S): Pharmacia Corporation, USA
 SOURCE: PCT Int. Appl., 266 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003088958	A2	20031030	WO 2003-US11269	
20030414 <--				
WO 2003088958	A3	20040819		
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RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
CA 2481934	A1	20031030	CA 2003-2481934	
20030414 <--				
AU 2003223579	A1	20031103	AU 2003-223579	
20030414 <--				
US 20040034083	A1	20040219	US 2003-413348	
20030414 <--				
EP 1494664	A2	20050112	EP 2003-719717	
20030414 <--				
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BR 2003009259	A	20050209	BR 2003-9259	
20030414 <--				
JP 2005528403	T	20050922	JP 2003-585710	
20030414 <--				
MX 2004009352	A	20050125	MX 2004-9352	
20040924 <--				
PRIORITY APPLN. INFO.:			US 2002-373311P	P
20020418 <--				
			WO 2003-US11269	W
20030414				

E KASE HIROSHI?/AU

L14	236 S E38
L15	15 S L14 AND L8
L16	15 S L15 AND (PY<2004 OR AY<2004 OR PRY<2004)
L17	14 S L16 NOT L10
L18	611 S SELECTIVE SEROTONIN REUPTAKE INHIBITORS/IT
L19	0 S L18 AND L8
L20	17 S L18 AND L3

L21 10 S L20 AND DEPRESSION?
 L22 4 S L21 AND (PY<2004 OR AY<2004 OR PRY<2004)

L22 ANSWER 1 OF 4 HCAPLUS COPYRIGHT 2009 ACS on STN
 TI Modified-release formulations of selective serotonin reuptake inhibitors
 ACCESSION NUMBER: 2009:769895 HCAPLUS Full-text
 DOCUMENT NUMBER: 151:86724
 TITLE: Modified-release formulations of selective serotonin reuptake inhibitors
 INVENTOR(S): Maes, Paul J.; Muhuri, Goutam
 PATENT ASSIGNEE(S): Biovail Laboratories Incorporated, Barbados
 SOURCE: Can., 178pp.
 CODEN: CAXXA4
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 2
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
CA 2415154	C	20090616	CA 2002-2415154	
20021224 <--				
CA 2415154	A1	20040624		
WO 2004058229	A1	20040715	WO 2003-CA1986	
20031219 <--				
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RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
AU 2003292927	A1	20040722	AU 2003-292927	
20031219 <--				
EP 1633329	A1	20060315	EP 2003-788728	
20031219 <--				
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US 20080138411	A1	20080612	US 2007-556492	
20070606 <--				
PRIORITY APPLN. INFO.:			CA 2002-2415154	A

20021224 <--

WO 2003-CA1986 W

20031219 <--

L22 ANSWER 2 OF 4 HCAPLUS COPYRIGHT 2009 ACS on STN

TI Modified-release formulations of selective serotonin reuptake inhibitors

ACCESSION NUMBER: 2004:565069 HCAPLUS Full-text

DOCUMENT NUMBER: 141:111573

TITLE: Modified-release formulations of selective serotonin

reuptake inhibitors

INVENTOR(S): Maes, Paul Jose; Muhuri, Goutam

PATENT ASSIGNEE(S): Biovail Laboratories Inc., Barbados

SOURCE: PCT Int. Appl., 214 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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WO 2004058229	A1	20040715	WO 2003-CA1986	
20031219 <--				
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GB, GD,				
GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR,				
KZ, LC,				
LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ,				
NI, NO,				
NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL,				
SY, TJ,				
TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW,				
AM, AZ,				
BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE,				
DK, EE,				
ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE,				
SI, SK,				
TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE,				
SN, TD, TG				
CA 2415154	C	20090616	CA 2002-2415154	
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CA 2415154	A1	20040624		
AU 2003292927	A1	20040722	AU 2003-292927	
20031219 <--				
EP 1633329	A1	20060315	EP 2003-788728	
20031219 <--				
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE,				
MC, PT,				
IE, SI, FI, RO, CY, TR, BG, CZ, EE, HU, SK				

L22 ANSWER 3 OF 4 HCAPLUS COPYRIGHT 2009 ACS on STN
TI Treatment of antidepressant drug-induced sexual dysfunction with
apomorphine

	713,741, abandoned.
	CODEN: USXXCO
DOCUMENT TYPE:	Patent
LANGUAGE:	English
FAMILY ACC. NUM. COUNT:	1
PATENT INFORMATION:	

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 20020086876 20011010 <-- PRIORITY APPLN. INFO.: 20001115 <-- IC ICM A61K031-4748 INCL 514295000 CC 1-12 (Pharmacology) IT Mental and behavioral disorders (depression; apomorphine for treatment of antidepressant-induced sexual dysfunction)	A1	20020704	US 2001-974136 US 2000-713741	B1
IT Biological transport (uptake, selective serotonin reuptake inhibitors; apomorphine for treatment of antidepressant-induced sexual dysfunction)				
IT 56296-78-7, Prozac 78246-49-8, Paxil 79559-97-0, Zoloft RL: ADV (Adverse effect, including toxicity); PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (apomorphine for treatment of antidepressant-induced sexual dysfunction, and use with antiemetics)				
OS.CITING REF COUNT:	1	THERE ARE 1 CAPLUS RECORDS THAT CITE THIS RECORD		
		(1 CITINGS)		

Ralph;
 Rowena;
 Weinberger,
 Gilsenan, Alicia; Eckert, George J.; Dolor,
 Stang, Paul; Zhou, Xiao-Hua; Hays, Ron;
 Morris
 CORPORATE SOURCE: Regenstrief Inst. Health Care, Indianapolis,
 IN,
 46202, USA
 SOURCE: JAMA, the Journal of the American Medical
 Association
 (2001), 286(23), 2947-2955
 CODEN: JAMAAP; ISSN: 0098-7484
 PUBLISHER: American Medical Association
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 CC 1-11 (Pharmacology)
 IT Antidepressants
 (selective serotonin reuptake
 inhibitors (SSRIs); similar effectiveness of paroxetine,
 fluoxetine, and sertraline in primary care)
 IT 56296-78-7, Fluoxetine hydrochloride 61869-08-7, Paroxetine
 79617-96-2, Sertraline
 RL: ADV (Adverse effect, including toxicity); PAC (Pharmacological
 activity); THU (Therapeutic use); BIOL (Biological study); USES
 (Uses)
 (similar effectiveness of paroxetine, fluoxetine, and
 sertraline in
 primary care)

L23 4 S L8 AND DEPRESSION?
 L24 3 S L23 AND (PY<2004 OR AY<2004 OR PRY<2004)

L24 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2009 ACS on STN
 TI Adenosine A2a receptor antagonists for the treatment of
 extrapyramidal
 syndrome and other movement disorders
 ACCESSION NUMBER: 2006:463565 HCAPLUS Full-text
 DOCUMENT NUMBER: 144:460860
 TITLE: Adenosine A2a receptor antagonists for the
 treatment
 of extrapyramidal syndrome and other movement
 disorders
 INVENTOR(S): Grzelak, Michael; Hunter, John; Pond,
 Annamarie;
 Varty, Geoffrey
 PATENT ASSIGNEE(S): Schering Corp., USA
 SOURCE: U.S. Pat. Appl. Publ., 27 pp., Cont.-in-part
 of U.S.
 Ser. No. 738,906.
 CODEN: USXXCO
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 3
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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US 20060106040	A1	20060518	US 2005-234644	
20050923 <--				
US 20040138235	A1	20040715	US 2003-738906	
20031217 <--				
US 7414058	B2	20080819		
CA 2510655	A1	20050519	CA 2003-2510655	
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AU 2003304527	A1	20050526	AU 2003-304527	
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EP 1578409	A1	20050928	EP 2003-818838	
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IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
BR 2003017436	A	20051116	BR 2003-17436	
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JP 2006514697	T	20060511	JP 2005-510511	
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NZ 540493	A	20080430	NZ 2003-540493	
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CN 101310724	A	20081126	CN 2008-10136154	
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MX 2005006790	A	20050908	MX 2005-6790	
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US 20060128694	A1	20060615	US 2005-249796	
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AU 2006294919	A1	20070405	AU 2006-294919	
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CA 2623040	A1	20070405	CA 2006-2623040	
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WO 2007038212	A1	20070405	WO 2006-US36864	
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GB, GD,				
GE, GH, GM, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM,				
KN, KP,				
MK, MN,				
MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT,				
RO, RS,				
RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR,				
TT, TZ,				
UA, UG, US, UZ, VC, VN, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR,				
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IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR,				
BF, BJ,				
CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG,				
BW, GH,				
GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM,				
AZ, BY,				

KG, KZ, MD, RU, TJ, TM
 EP 1940408 A1 20080709 EP 2006-815125
 20060921
 R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR,
 HU, IE,
 IS, IT, LI, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK,
 TR, AL,
 BA, HR, MK, RS
 JP 2009508967 T 20090305 JP 2008-532393
 20060921
 ZA 2008002552 A 20090624 ZA 2008-2552
 20080319
 MX 2008004006 A 20080410 MX 2008-4006
 20080324
 CN 101312731 A 20081126 CN 2006-80043146
 20080519
 PRIORITY APPLN. INFO.: US 2002-435321P P
 20021219 <--
 US 2003-738906 A2
 20031217 <--
 CN 2003-80107087 A3
 20031217 <--
 WO 2003-US40456 W
 20031217 <--
 US 2005-234644 A2
 20050923
 WO 2006-US36864 W
 20060921
 OTHER SOURCE(S): MARPAT 144:460860
 INCL 514263240; 514263340
 CC 1-11 (Pharmacology)
 IT Mental and behavioral disorders
 (depression, dystonia from antidepressant use; adenosine A2a
 receptor antagonists for treatment of extrapyramidal syndrome
 and other
 movement disorders)
 IT 59-92-7, Levodopa, biological studies 322-35-0, Benserazide
 7439-89-6D, Iron, salts 12794-10-4D, Benzodiazepine, derivs.
 28860-95-9, Carbidopa 155270-99-8 377727-26-9
 RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL
 (Biological study); USES (Uses)
 (adenosine A2a receptor antagonists for treatment of
 extrapyramidal
 syndrome and other movement disorders)
 L24 ANSWER 2 OF 3 HCAPLUS COPYRIGHT 2009 ACS on STN
 TI Medicinal compositions containing adenosine A2A receptor
 antagonists and
 other antidepressants
 ACCESSION NUMBER: 2005:99358 HCAPLUS Full-text
 DOCUMENT NUMBER: 142:162694
 TITLE: Medicinal compositions containing adenosine
 A2A
 receptor antagonists and other antidepressants
 INVENTOR(S): Kase, Hiroshi; Kobayashi, Minoru; Shiozaki,
 Shizuo;
 Mori, Akihisa; Seno, Naoki

PATENT ASSIGNEE(S): Kyowa Hakko Kogyo Co., Ltd., Japan
 SOURCE: PCT Int. Appl., 47 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: Japanese
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005009444	A1	20050203	WO 2004-JP10758	
20040722 <--				
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
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CA 2533117	A1	20050203	CA 2004-2533117	
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EP 1655029	A1	20060510	EP 2004-748023	
20040722 <--				
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US 20060241102	A1	20061026	US 2006-565239	
20060119 <--				
NO 2006000958	A	20060425	NO 2006-958	
20060227 <--				
PRIORITY APPLN. INFO.:			JP 2003-201549	A
20030725 <--			WO 2004-JP10758	W
20040722				
IC ICM A61K031-52				
ICS A61K031-137; A61K031-335; A61K031-343; A61K031-36; A61K031-38;				
A61K031-381; A61K031-435; A61K031-496; A61K031-5375; A61K031-55;				
A61K031-553; A61P025-24				
CC 63-6 (Pharmaceuticals)				
Section cross-reference(s): 1				

IT Mental and behavioral disorders
 (depression; medicinal compns. containing adenosine A2A
 receptor
 antagonists and other antidepressants)
 IT 56296-78-7, Fluoxetine hydrochloride 99300-78-4, Venlafaxine
 hydrochloride 155270-99-8
 RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL
 (Biological study); USES (Uses)
 (medicinal compns. containing adenosine A2A receptor
 antagonists and other
 antidepressants)
 OS.CITING REF COUNT: 3 THERE ARE 3 CAPLUS RECORDS THAT CITE
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 (3 CITINGS)
 REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE
 FOR THIS
 RECORD. ALL CITATIONS AVAILABLE IN THE
 RE FORMAT

L24 ANSWER 3 OF 3 HCAPLUS COPYRIGHT 2009 ACS on STN
 TI KW-6002, Kyowa Hakko Kogyo
 ACCESSION NUMBER: 2001:438063 HCAPLUS Full-text
 DOCUMENT NUMBER: 136:193429
 TITLE: KW-6002, Kyowa Hakko Kogyo
 AUTHOR(S): Knutsen, Lars J. S.; Weiss, Scott M.
 CORPORATE SOURCE: Vernalis Research Limited, Berkshire, RG41
 5UA, UK
 SOURCE: Current Opinion in Investigational Drugs
 (PharmaPress
 Ltd.) (2001), 2(5), 668-673
 CODEN: COIDAZ
 PUBLISHER: PharmaPress Ltd.
 DOCUMENT TYPE: Journal; General Review
 LANGUAGE: English
 CC 1-0 (Pharmacology)
 IT 155270-99-8, KW-6002
 RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL
 (Biological study); USES (Uses)
 (KW-6002 for treatment of Parkinson's disease)
 OS.CITING REF COUNT: 22 THERE ARE 22 CAPLUS RECORDS THAT CITE
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 RECORD (22 CITINGS)
 REFERENCE COUNT: 38 THERE ARE 38 CITED REFERENCES AVAILABLE
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 RECORD. ALL CITATIONS AVAILABLE IN THE
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=> d 124 ti abs 3

L24 ANSWER 3 OF 3 HCAPLUS COPYRIGHT 2009 ACS on STN
 TI KW-6002, Kyowa Hakko Kogyo
 AB A review. Kyowa Hakko is developing KW-6002, an adenosine A2A
 receptor antagonist, for the potential treatment of Parkinson's
 disease (PD). The company subsequently began evaluating the
 compound for depression, and in June 2000 initiated a phase II
 trial for this indication. Good results have been obtained in a

preclin. depression model and Kyowa Hakko hopes that KW-6002 will be representative of a new class of antidepressants. KW-6002 is undergoing phase II trials for PD in Japan and in Europe, including the UK. The drug is of particular interest due to the absence of the involuntary movement adverse effects characteristic of L-DOPA therapy. Roche was to codevelop KW-6002 with Kyowa in the US and Europe, but withdrew from the development of this drug in Apr. 1999. Kyowa will continue with the drug's development in the US. NDAs are estimated to be filed around the end of 2001 and the drug is expected to be launched in Western Europe during 2002. An NDA is expected to be filed in Japan approx. three to four years later than in Western Europe.

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E KOBAYASHI MINORU?/AU
L25      401 S E49-E50
L26      4 S L25 AND L8
L27      3 S L26 NOT L10
L28      3 S L27 NOT L24
E SHIOZAKI SHIZUO?/AU
L29      49 S E61-E62
L30      8 S L29 AND L8
L31      7 S L30 NOT L10
L32      7 S L31 NOT L24
L33      7 S L32 AND (PY<2004 OR AY<2004 OR PRY<2004)
E MORI AKIHISA?/AU
L34      187 S E73-E74
L35      6 S L34 AND (L8 OR L3)
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L37      4 S L36 AND (PY<2004 OR AY<2004 OR PRY<2004)
L38      3 S L37 NOT L33
E SENO NAOKI?/AU
L39      37 S E85-E86
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L41      1 S L40 NOT (L10 OR L33)

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